

Notice of References Cited	Application/Control No. 10/021,955	Applicant(s)/Patent Under Reexamination LUPSKI ET AL.	
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U.S. PATENT DOCUMENTS

*		Document Number Country Code-Number-Kind Code	Date MM-YYYY	Name	Classification
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*		Include as applicable: Author, Title Date, Publisher, Edition or Volume, Pertinent Pages)
*	U	Kijima et al. Periaxin mutation causes early-onset but slow-progressive Charcot-Marie-Tooth disease. J Hum Genet., Vol. 49, pp. 376-379, 2004.
*	V	Takashima et al. Periaxin mutations cause a broad spectrum of demyelinating neuropathies. Ann NEurol., Vol. 51, pp. 709-715 2002.
*	W	Gillespie et al. The gene encoding the schwann cell protein periaxin localizes on mouse chromosome 7 (PRX). Geneomics, Vo 41, pages 297-298, 1997.
*	X	Boerkoel et al. Periaxin mutations cause recessive Dejerine-Sottas Neuropathy. Am. J. Hum. Genet., Vol. 68, pages 325-333, 2001.

*A copy of this reference is not being furnished with this Office action. (See MPEP § 707.05(a).)
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*	U	Timmerman et al. Novel missense mutation in the early growth response 2 gene associated with Dejerine-Sttas syndrome phenotype. <i>Neurology</i> , Vol. 52, pp. 1827-1832, 1999.
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